## **AMENDMENTS TO THE CLAIMS**

- 1 43. (Canceled).
- 44. (Previously presented) A method for inhibiting immunoglobulin production comprising contacting T-cells with an antibody that specifically binds to a protein specifically recognized by monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048.
- 45. (Canceled)
- 46. (Previously presented) A method for inhibiting activation of B-cells comprising contacting T-cells with an antibody that specifically binds to a protein specifically recognized by monoclonal antibody MRI produced by the hybridoma having ATCC Accession No. HB 11048.
- 47. (Canceled)
- 48. (Canceled)
- 49. (Canceled)
- 50. (Currently amended) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal, in an amount effective to inhibit immunoglobulin production, an antibody that specifically binds to a protein specifically recognized by the monoclonal antibody MRl produced by the hybridoma having ATCC Accession No. HB 11048.
- 51. (Canceled)
- 52. (Previously presented) A method for inhibiting activation of B-cells in an animal comprising administering to the animal, in an amount effective to inhibit activation of B-cells, an antibody that specifically binds to a protein

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specifically recognized by monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048.

## 53. (Canceled)

- 54. (Currently amended) The method of any one of claims 42 44, 46, 50, and 52 through 53, wherein the antibody is selected from the group consisting of monoclonal antibodies, chimeric antibodies, human antibodies, and fragments thereof that specifically bind to a protein specifically bound by the MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048.
- 55. (Currently amended) The method of any of claims 42 through 53 44, 46, 50, and 52, wherein the antibody further comprises is conjugated to a moiety selected from the group consisting of an enzyme, a toxin, a growth factor, a lymphokine, an anti-proliferative agent, an alkylating agent, an antimetabolite, an antibiotic, a vinca alkaloid, a platinum coordinated complex, a radioisotope, and a fluorescent compound, wherein the moiety is conjugated to the antibody.
- 56. (Currently amended) The method of any one of claims 42 through 53 44, 46, 50, and 52, wherein the antibody is conjugated to further comprises a therapeutic agent, wherein the therapeutic agent is conjugated to the antibody.
- 57. (Currently amended) The method of any of claims 48 through 53 50 and 52, wherein the animal is a mammal.
- 58. (Currently amended) The method of any of claims 48 through 53 50 and 52, wherein the animal is a human.
- 59. (Canceled)

60. (Previously presented; allowed) A method for inhibiting immunoglobulin production comprising contacting T-cells with a composition comprising monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048, and fragments thereof that specifically bind to a protein specifically bound by the MR1 antibody.

- 61. (Previously presented; allowed) A method for inhibiting activation of B-cells comprising contacting T-cells with a composition comprising monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048, and fragments thereof that specifically bind to a protein specifically bound by the MR1 antibody.
- 62. (Canceled).
- 63. (Previously presented; allowed) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal, in an amount effective to inhibit immunoglobulin production, a composition comprising monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048, and fragments thereof that specifically bind to a protein specifically bound by the MR1 antibody.
- 64. (Previously presented; allowed) A method for inhibiting activation of B-cells in an animal comprising administering to the animal, in an amount effective to inhibit activation of B-cells, a composition comprising monoclonal antibody MR1 produced by the hybridoma having ATCC Accession No. HB 11048, and fragments thereof that specifically bind to a protein specifically bound by the MR1 antibody.
- 65. (Canceled)

66. (Previously presented; allowed) A method for inhibiting immunoglobulin production comprising contacting T-cells with a human antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.

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- 67. (Previously presented; allowed) A method for inhibiting activation of B-cells comprising contacting T-cells with a human antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.
- 68. (Canceled)
- 69. (Previously presented; allowed) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal, in an amount effective to inhibit immunoglobulin production, a composition comprising a human antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.
- 70. (Previously presented; allowed) A method for inhibiting activation of B-cells in an animal comprising administering to the animal, in an amount effective to inhibit activation of B-cells, a composition comprising a human antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.

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## 71. (Canceled)

- 72. (Previously presented; allowed) A method for inhibiting immunoglobulin production comprising contacting T-cells with a chimeric antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.
- 73. (Previously presented; allowed) A method for inhibiting activation of B-cells comprising contacting T-cells with a chimeric antibody comprising a binding fragment of monoclonal MRI antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MRI antibody.
- 74. (Canceled).
- 75. (Previously presented; allowed) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal, in an amount effective to inhibit immunoglobulin production, a composition comprising a chimeric antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein the binding fragment specifically binds to a protein specifically bound by the MR1 antibody.
- 76. (Previously presented; allowed) A method for inhibiting activation of B-cells in an animal comprising administering to the animal, in an amount effective to inhibit activation of B-cells, a composition comprising a chimeric antibody comprising a binding fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048, wherein

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the binding fragment specifically binds to a protein specifically bound by the MRl antibody.

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- 77. (Canceled)
- 78. (Previously presented; allowed) A method for inhibiting immunoglobulin production comprising contacting T-cells with an F(ab')<sub>2</sub> fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048.
- 79. (Previously presented; allowed) A method for inhibiting activation of B-cells comprising contacting T-cells with a chimeric antibody comprising an F(ab')<sub>2</sub> fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048.
- 80. (Canceled).
- 81. (Previously presented; allowed) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal, in an amount effective to inhibit immunoglobulin production, a composition comprising an F(ab')<sub>2</sub> fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048.
- 82. (Previously presented; allowed) A method for inhibiting activation of B-cells in an animal comprising administering to the animal, in an amount effective to inhibit activation of B-cells, a composition comprising an F(ab')<sub>2</sub> fragment of monoclonal MR1 antibody produced by the hybridoma having ATCC Accession No. HB 11048.

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83. (New) A method for inhibiting immunoglobulin production comprising contacting T-cells with an effective amount of an antibody that binds an antigen that:

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- (a) is present on activated but not resting T-cells;
- (b) has the same molecular weight as a protein precipitated by a CD40-immunoglobulin fusion protein (CD40-Ig), the CD40-Ig comprising the extracellular domain of a CD40 protein having the amino acid sequence of SEQ ID NO:2 and an extracellular domain at the site of fusion having the amino acid sequence of SEQ ID NO:3; and
- (c) is pre-cleared by precipitation with the CD40-Ig; wherein the antibody blocks binding of the CD40-Ig to activated T-cells and inhibits T-cell induction of B-cell activation.
- 84. (New) A method for inhibiting activation of B-cells comprising contacting T-cells with an effective amount of an antibody that binds an antigen that:
  - (a) is present on activated but not resting T-cells;
- (b) has the same molecular weight as a protein precipitated by a CD40-immunoglobulin fusion protein (CD40-Ig), the CD40-Ig comprising the extracellular domain of a CD40 protein having the amino acid sequence of SEQ ID NO:2 and an extracellular domain at the site of fusion having the amino acid sequence of SEQ ID NO:3; and
- (c) is pre-cleared by precipitation with the CD40-Ig; wherein the antibody blocks binding of the CD40-Ig to activated T-cells and inhibits T-cell induction of B-cell activation.
- 85. (New) A method for inhibiting immunoglobulin production in an animal comprising the step of administering to the animal an effective amount of an antibody that binds an antigen that:
  - (a) is present on activated but not resting T-cells;

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- (b) has the same molecular weight as a protein precipitated by a CD40-immunoglobulin fusion protein (CD40-Ig), the CD40-Ig comprising the extracellular domain of a CD40 protein having the amino acid sequence of SEQ ID NO:2 and an extracellular domain at the site of fusion having the amino acid sequence of SEQ ID NO:3; and
- (c) is pre-cleared by precipitation with the CD40-Ig; wherein the antibody blocks binding of the CD40-Ig to activated T-cells and inhibits T-cell induction of B-cell activation.
- 86. (New) A method for inhibiting activation of B-cells in an animal comprising administering to the animal an effective amount of an antibody that binds an antigen that:
  - (a) is present on activated but not resting T-cells;
- (b) has the same molecular weight as a protein precipitated by a CD40-immunoglobulin fusion protein (CD40-Ig), the CD40-Ig comprising the extracellular domain of a CD40 protein having the amino acid sequence of SEQ ID NO:2 and an extracellular domain at the site of fusion having the amino acid sequence of SEQ ID NO:3; and
- (c) is pre-cleared by precipitation with the CD40-Ig; wherein the antibody blocks binding of the CD40-Ig to activated Tcells and inhibits T-cell induction of B-cell activation.
- 87. (New) The method of any of claims 83-86, wherein the antibody is conjugated to a moiety selected from the group consisting of an enzyme, a toxin, a growth factor, a lymphokine, an anti-proliferative agent, an alkylating agent, an anti-metabolite, an antibiotic, a vinca alkaloid, a platinum coordinated complex, a radioisotope, and a fluorescent compound.
- 88. (New) The method of any one of claims 83-86, wherein the antibody is conjugated to a therapeutic agent.

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89. (New) The method of any of claims 85 and 86, wherein the animal is a mammal.

90. (New) The method of any of claims 85 and 86, wherein the animal is a mouse.